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(54) Expandable medical device for delivery of beneficial agent

(57) The invention provides an expandable medical device (100), comprising: a plurality of elongated struts (18) forming a substantially cylindrical device (12) which is expandable from a cylinder having a first diameter to a cylinder having a second diameter; hinges (20) formed

at said elongated struts (18) to substantially prevent deformation of said elongated struts (18) during expansion of said cylindrical device (12), and links (22) arranged between said elongated struts (18); wherein a series of openings (24, 26) are formed in said struts (18) and/or said links (22).

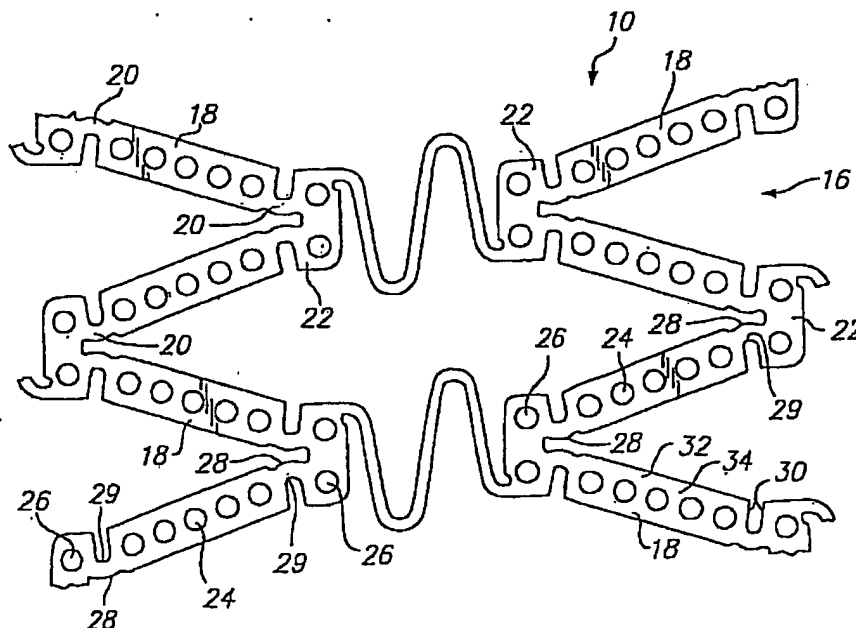


FIG. 2

EP 1 582 180 A1

Description

BACKGROUND OF THE INVENTION

Field of the Invention

[0001] The present invention relates to tissue-supporting medical devices, and more particularly to expandable, non-removable devices that are implanted within a bodily lumen of a living animal or human to support the organ and maintain patency, and that can deliver a beneficial agent to the intervention site.

Summary of the Related Art

[0002] In the past, permanent or biodegradable devices have been developed for implantation within a body passageway to maintain patency of the passageway. These devices are typically introduced percutaneously, and transported transluminally until positioned at a desired location. These devices are then expanded either mechanically, such as by the expansion of a mandrel or balloon positioned inside the device, or expand themselves by releasing stored energy upon actuation within the body. Once expanded within the lumen, these devices, called stents, become encapsulated within the body tissue and remain a permanent implant.

[0003] Known stent designs include monofilament wire coil stents (U.S. Pat. No. 4,969,458); welded metal cages (U.S. Pat. Nos. 4,733,665 and 4,776,337); and, most prominently, thin-walled metal cylinders with axial slots formed around the circumference (U.S. Pat. Nos. 4,733,665, 4,730,762, and 4,776,337). Known construction materials for use in stents include polymers, organic fabrics and biocompatible metals, such as, stainless steel, gold, silver, tantalum, titanium, and shape memory alloys such as Nitinol.

[0004] U.S. Pat. Nos. 4,733,665, 4,739,762, and 4,776,337 disclose expandable and deformable interluminal vascular grafts in the form of thin-walled tubular members with axial slots allowing the members to be expanded radially outwardly into contact with a body passageway. After insertion, the tubular members are mechanically expanded beyond their elastic limit and thus permanently fixed within the body. The force required to expand these tubular stents is proportional to the thickness of the wall material in a radial direction. To keep expansion forces within acceptable levels for use within the body (e.g., 5 - 10 atm), these designs must use very thin-walled materials (e.g., stainless steel tubing with 0.0025 inch thick walls). However, materials this thin are not visible on conventional fluoroscopic and x-ray equipment and it is therefore difficult to place the stents accurately or to find and retrieve stents that subsequently become dislodged and lost in the circulatory system.

[0005] Further, many of these thin-walled tubular stent designs employ networks of long, slender struts

whose width in a circumferential direction is two or more times greater than their thickness in a radial direction. When expanded, these struts are frequently unstable, that is, they display a tendency to buckle, with individual struts twisting out of plane. Excessive protrusion of these twisted struts into the bloodstream has been observed to increase turbulence, and thus encourage thrombosis. Additional procedures have often been required to attempt to correct this problem of buckled struts. For example, after initial stent implantation is determined to have caused buckling of struts, a second, high-pressure balloon (e.g., 12 to 18 atm) would be used to attempt to drive the twisted struts further into the lumen wall. These secondary procedures can be dangerous to the patient due to the risk of collateral damage to the lumen wall.

[0006] Many of the known stents display a large elastic recovery, known in the field as "recoil," after expansion inside a lumen. Large recoil necessitates over-expansion of the stent during implantation to achieve the desired final diameter. Over-expansion is potentially destructive to the lumen tissue. Known stents of the type described above experience recoil of up to about 6 to 12% from maximum expansion.

[0007] Large recoil also makes it very difficult to securely crimp most known stents onto delivery catheter balloons. As a result, slippage of stents on balloons during interluminal transportation, final positioning, and implantation has been an ongoing problem. Many ancillary stent securing devices and techniques have been advanced to attempt to compensate for this basic design problem. Some of the stent securing devices include collars and sleeves used to secure the stent onto the balloon.

[0008] Another problem with known stent designs is non-uniformity in the geometry of the expanded stent. Non-uniform expansion can lead to non-uniform coverage of the lumen wall creating gaps in coverage and inadequate lumen support. Further, over expansion in some regions or cells of the stent can lead to excessive material strain and even failure of stent features. This problem is potentially worse in low expansion force stents having smaller feature widths and thicknesses in which manufacturing variations become proportionately more significant. In addition, a typical delivery catheter for use in expanding a stent includes a balloon folded into a compact shape for catheter insertion. The balloon is expanded by fluid pressure to unfold the balloon and deploy the stent. This process of unfolding the balloon causes uneven stresses to be applied to the stent during expansion of the balloon due to the folds causing the problem non-uniform stent expansion.

[0009] U.S. Pat. No. 5,545,210 discloses a thin-walled tubular stent geometrically similar to those discussed above, but constructed of a nickel-titanium shape memory alloy ("Nitinol"). This design permits the use of cylinders with thicker walls by making use of the lower yield stress and lower elastic modulus of marten-

silic phase Nitinol alloys. The expansion force required to expand a Nitinol stent is less than that of comparable thickness stainless steel stents of a conventional design. However, the "recoil" problem after expansion is significantly greater with Nitinol than with other materials. For example, recoil of a typical design Nitinol stent is about 9%. Nitinol is also more expensive, and more difficult to fabricate and machine than other stent materials, such as stainless steel.

[0010] All of the above stents share a critical design property: in each design, the features that undergo permanent deformation during stent expansion are prismatic, i.e., the cross sections of these features remain constant or change very gradually along their entire active length. To a first approximation, such features deform under transverse stress as simple beams with fixed or guided ends: essentially, the features act as a leaf springs. These leaf spring like structures are ideally suited to providing large amounts of elastic deformation before permanent deformation commences. This is exactly the opposite of ideal stent behavior. Further, the force required to deflect prismatic stent struts in the circumferential direction during stent expansion is proportional to the square of the width of the strut in the circumferential direction. Expansion forces thus increase rapidly with strut width in the above stent designs. Typical expansion pressures required to expand known stents are between about 5 and 10 atmospheres. These forces can cause substantial damage to tissue if misapplied.

[0011] In addition to the above-mentioned risks to a patient, restenosis is a major complication which can arise following the implantation of stents, using stent devices such as those described above, and other vascular interventions such as angioplasty. Simply defined, restenosis is a wound healing process that reduces the vessel lumen diameter by scar tissue formation and which may ultimately result in reocclusion of the lumen. Despite the introduction of improved surgical techniques, devices and pharmaceutical agents, the overall restenosis rate is still reported in the range of 25% to 50% within six to twelve months after an angioplasty procedure. To correct this problem, additional revascularization procedures are frequently required, thereby increasing trauma and risk to the patient.

[0012] Several techniques under development to address the problem of restenosis are irradiation of the injury site and the use of stents to deliver a variety of beneficial or pharmaceutical agents to the traumatized vessel lumen. In the latter case, a stent is frequently surface-coated with a beneficial agent (often a drug-impregnated polymer) and implanted at the angioplasty site. Alternatively, an external drug-impregnated polymer sheath is mounted over the stent and co-deployed in the vessel. In either case, it has proven difficult to deliver a sufficient amount of beneficial agent to the trauma site so as to satisfactorily prevent the growth of scar tissue and thereby reduce the likelihood of restenosis. Even with relatively thick coatings of the beneficial agent

or sheaths of increased thickness surrounding the stents, restenosis has been found to occur. Furthermore, increasing the effective stent thickness (e.g., by providing increased coatings of the beneficial agent) is undesirable for a number of reasons, including increased trauma to the vessel lumen during implantation and reduced flow cross-section of the lumen after implantation. Moreover, coating thickness is one of several factors that affect the release kinetics of the beneficial agent, and limitations on thickness thereby limit the range of release rates, durations, and the like that can be achieved.

SUMMARY OF THE INVENTION

[0013] In view of the drawbacks of the prior art, it would be advantageous to provide a stent capable of delivering a relatively large volume of a beneficial agent to a traumatized site in a vessel lumen without increasing the effective wall thickness of the stent, and without adversely impacting the mechanical expansion properties of the stent.

[0014] It would further be advantageous to have such a stent, which also significantly increases the available depth of the beneficial agent reservoir.

[0015] It would be further advantageous to be able to expand such a stent with an expansion force at a low level independent of choice of stent materials, material thickness, or strut dimensions.

[0016] It would further be advantageous to have such a tissue-supporting device that permits a choice of material thickness that could be viewed easily on conventional fluoroscopic equipment for any material.

[0017] It would also be advantageous to have such a tissue-supporting device that is inherently stable during expansion, thus eliminating buckling and twisting of structural features during stent deployment.

[0018] In addition, it would be advantageous to have such a tissue-supporting device with minimal elastic recovery, or "recoil" of the device after expansion.

[0019] It would be advantageous to have such a tissue supporting device that can be securely crimped to the delivery catheter without requiring special tools, techniques, or ancillary clamping features.

[0020] In accordance with one aspect of the invention, an expandable medical device includes a cylindrical tube, and a network of elongated struts formed in the cylindrical tube, wherein each of the elongated struts are axially displaced from adjacent struts. A plurality of ductile hinges are formed between the elongated struts. The ductile hinges allow the cylindrical tube to be expanded or compressed from a first diameter to a second diameter by deformation of the ductile hinges. Further, at least one of the elongated struts includes at least one opening for loading of a beneficial agent therein. The at least one opening may include a plurality of openings that extend through a thickness of the at least one strut, so as to thereby define a through-opening, or

the openings may have a depth less than a thickness of the at least one strut, so as to thereby define a recess. A beneficial agent is loaded within the at least one opening, wherein the beneficial agent includes antiproliferatives, antithrombins, large molecules, microspheres, biodegradable agents, or cells. The at least one opening of the at least one strut forms a protected receptor for loading the beneficial agent therein.

[0021] In accordance with a further aspect of the present invention, an expandable medical device includes a plurality of elongated struts, the plurality of elongated struts joined together to form a substantially cylindrical device which is expandable from a cylinder having a first diameter to a cylinder having a second diameter, and the plurality of struts each having a strut width in a circumferential direction. At least one of the plurality of struts includes at least one recess extending at least partially through a thickness of the strut. The at least one recess may extend entirely through the thickness of the strut so as to define a through-opening and the at least one recess may be generally rectangular or polygonal.

BRIEF DESCRIPTION OF THE DRAWINGS

[0022] The invention will now be described in greater detail with reference to the preferred embodiments illustrated in the accompanying drawings, in which like elements bear like reference numerals, and wherein:

FIG. 1 is a perspective view of a tissue-supporting device in accordance with a first preferred embodiment of the present invention;

FIG. 2 is an enlarged side view of a portion thereof;

FIG. 3 is an enlarged side view of a tissue-supporting device in accordance with a further preferred embodiment of the present invention;

FIG. 4 is an enlarged side view of a portion of the stent shown in the device of FIG. 3;

FIG. 5 is an enlarged cross section of an opening thereof;

FIG. 6 is an enlarged cross section of an opening thereof illustrating beneficial agent loaded into the opening;

FIG. 7 is an enlarged cross section of an opening thereof illustrating a beneficial agent loaded into the opening and a thin coating of a beneficial agent;

FIG. 8 is an enlarged cross section of an opening thereof illustrating a beneficial agent loaded into the opening and thin coatings of different beneficial agents on different surfaces of the device;

FIG. 9 is an enlarged side view of a portion of a stent in accordance with yet another preferred embodiment of the present invention;

FIGS. 10a - 10c are perspective, side, and cross-sectional views of an idealized ductile hinge for purposes of analysis, and FIG. 10d is a stress/strain curve for the idealized ductile hinge;

FIGS. 11 is a perspective view of a simple beam for purposes of calculation;

FIG. 12 is a moment verses curvature graph for a rectangular beam; and

FIG. 13 is an enlarged side view of a bent ductile hinge.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0023] Referring to FIGS. 1 and 2, a tissue supporting device in accordance with a preferred embodiment of the present invention is shown generally by reference numeral 10. The tissue supporting device 10 includes a plurality of cylindrical tubes 12 connected by S-shaped bridging elements 14. The bridging elements 14 allow the tissue supporting device to bend axially when passing through the tortuous path of the vasculature to the deployment site and allow the device to bend when necessary to match the curvature of a lumen to be supported. The S-shaped bridging elements 14 provide improved axial flexibility over prior art devices due to the thickness of the elements in the radial direction which allows the width of the elements to be relatively small without sacrificing radial strength. For example, the width of the bridging elements 14 may be about 0.0015 - 0.0018 inches (0.0381 - 0.0457 mm). Each of the cylindrical tubes 12 has a plurality of axial slots 16 extending from an end surface of the cylindrical tube toward an opposite end surface.

[0024] Formed between the slots 16 is a network of axial struts 18 and links 22. The cross section (and rectangular moment of inertia) of each of the struts 18 is preferably not constant along the length of the strut. Rather, the strut cross section changes abruptly at both ends of each strut 18 adjoining the links 22. The preferred struts 18 are thus not prismatic. Each individual strut 18 is preferably linked to the rest of the structure through a pair of reduced sections 20, one at each end, which act as stress/strain concentration features. The reduced sections 20 of the struts function as hinges in the cylindrical structure. Since the stress/strain concentration features are designed to operate into the plastic deformation range of generally ductile materials, they are referred to as ductile hinges 20. Such features are also commonly referred to as "Notch Hinges" or "Notch Springs" in ultra-precision mechanism design, where they are used exclusively in the elastic range.

[0025] With reference to the drawings and the discussion, the *width* of any feature is defined as its dimension in the circumferential direction of the cylinder. The *length* of any feature is defined as its dimension in the axial direction of the cylinder. The *thickness* of any feature is defined as the wall thickness of the cylinder.

[0026] Ductile hinges 20 are preferably asymmetric ductile hinges that produce different strain versus deflection-angle functions in expansion and compression. Each of the ductile hinges 20 is formed between a are

surface 28 and a concave notch surface 29. The ductile hinge 20 according to a preferred embodiment essentially takes the form of a small, prismatic curved beam having a substantially constant cross section. However, a thickness of the curved ductile hinge 20 may vary somewhat as long as the ductile hinge width remains constant along a portion of the hinge length. The width of the curved beam is measured along the radius of curvature of the beam. This small curved beam is oriented such that the smaller concave notch surface 29 is placed in tension in the device *crimping* direction, while the larger arc surface 28 of the ductile hinges is placed in tension in the device *expansion* direction. Again, there is no local minimum width of the ductile hinge 20 along the (curved) ductile hinge axis, and no concentration of material strain. During device expansion tensile strain will be distributed along the arc surface 28 of the hinge 20 and maximum expansion will be limited by the angle of the walls of the concave notch 29 which provide a geometric deflection limiting feature. The notches 29 each have two opposed angled walls 30 which function as a stop to limit geometric deflection of the ductile hinge, and thus limit maximum device expansion. As the cylindrical tubes 12 are expanded and bending occurs at the ductile hinges 20, the angled side walls 30 of the notches 29 move toward each other. Once the opposite side walls 30 of a notch come into contact with each other, they resist further expansion of the particular ductile hinge causing further expansion to occur at other sections of the tissue supporting device. This geometric deflection limiting feature is particularly useful where uneven expansion is caused by either variations in the tissue supporting device 10 due to manufacturing tolerances or uneven balloon expansion. Maximum tensile strain can therefore be reliably limited by adjusting the initial length of the arc shaped ductile hinge 20 over which the total elongation is distributed.

[0027] The presence of the ductile hinges 20 allows all of the remaining features in the tissue supporting device to be increased in width or the circumferentially oriented component of their respective rectangular moments of inertia - thus greatly increasing the strength and rigidity of these features. The net result is that elastic, and then plastic deformation commence and propagate in the ductile hinges 20 before other structural elements of the device undergo any significant elastic deformation. The force required to expand the tissue supporting device 10 becomes a function of the geometry of the ductile hinges 20, rather than the device structure as a whole, and arbitrarily small expansion forces can be specified by changing hinge geometry for virtually any material wall thickness. In particular, wall thicknesses great enough to be visible on a fluoroscope can be chosen for any material of interest.

[0028] In order to get minimum recoil, the ductile hinges 20 should be designed to operate well into the plastic range of the material, and relatively high local strain-curvatures are developed. When these conditions apply,

elastic curvature is a very small fraction of plastic or total curvature, and thus when expansion forces are relaxed, the percent change in hinge curvature is very small. When incorporated into a strut network designed to take maximum advantage of this effect, the elastic spring-back, or "recoil," of the overall stent structure is minimized.

[0029] In the preferred embodiment of FIGS. 1 and 2, it is desirable to increase the width of the individual struts 18 between the ductile hinges 20 to the maximum width that is geometrically possible for a given diameter and a given number of struts arrayed around that diameter. The only geometric limitation on strut width is the minimum practical width of the slots 16 which is about 0.002 inches (0.0508 mm) for laser machining. Lateral stiffness of the struts 18 increases as the cube of strut width, so that relatively small increases in strut width significantly increase strut stiffness. The net result of inserting ductile hinges 20 and increasing strut width is that the struts 18 no longer act as flexible leaf springs, but act as essentially rigid beams between the ductile hinges. All radial expansion or compression of the cylindrical tissue supporting device 10 is accommodated by mechanical strain in the hinge features 20, and yield in the hinge commences at very small overall radial expansion or compression.

[0030] Yield in ductile hinges at very low gross radial deflections also provides the superior crimping properties displayed by the ductile hinge-based designs. When a tissue supporting device is crimped onto a folded catheter balloon, very little radial compression of the device is possible since the initial fit between balloon and device is already snug. Most stents simply rebound elastically after such compression, resulting in very low clamping forces and the attendant tendency for the stent to slip on the balloon. Ductile hinges, however, sustain significant plastic deformation even at the low deflections occurring during crimping onto the balloon, and therefore a device employing ductile hinges displays much higher clamping forces. The ductile hinge designs according to the present invention may be securely crimped onto a balloon of a delivery catheter by hand or by machine without the need for auxiliary retaining devices commonly used to hold known stents in place.

[0031] The ductile hinge 20 illustrated in FIGS. 1 and 2 is exemplary of a preferred structure that will function as a stress/strain concentrator. Many other stress/strain concentrator configurations may also be used as the ductile hinges in the present invention, as shown and described for example in U.S. Application Serial No. 09/183,555, the entire contents of which is hereby incorporated by reference. The geometric details of the stress/strain concentration features or ductile hinges 20 can be varied greatly to tailor the exact mechanical expansion properties to those required in a specific application. The ductile hinges according to the present invention generally include an abrupt change in width of a strut that functions to concentrate stresses and strains

in the narrower section of the strut. These ductile hinges also generally include features to limit mechanical deflection of attached struts and features to control material strain during large strut deflections. Although the ductile hinges have been illustrated in FIG. 2 as positioned along the length of the struts 18 and the links 22, they may also be positioned at other locations in other designs without departing from the present invention.

[0032] At intervals along the neutral axis of the struts 18, at least one and more preferably a series of through-openings 24 are formed by laser drilling or any other means known to one skilled in the art. Similarly, at least one and preferably a series of through-openings 26 are formed at selected locations in the links 22. Although the use of through-openings 24 and 26 in both the struts 18 and links 22 is preferred, it should be clear to one skilled in the art that through-openings could be formed in only one of the struts and links. In the illustrated embodiment, the through-openings 24, 26 are circular in nature and thereby form cylindrical holes extending through the width of the tissue supporting device 10. It should be apparent to one skilled in the art, however, that through-openings of any geometrical shape or configuration could of course be used without departing from the scope of the present invention.

[0033] The behavior of the struts 18 in bending is analogous to the behavior of an I-beam or truss. The outer edge elements 32 of the struts 18 correspond to the I-beam flange and carry the tensile and compressive stresses, whereas the inner elements 34 of the struts 18 correspond to the web of an I-beam which carries the shear and helps to prevent buckling and wrinkling of the faces. Since most of the bending load is carried by the outer edge elements 32 of the struts 18, a concentration of as much material as possible away from the neutral axis results in the most efficient sections for resisting strut flexure. As a result, material can be judiciously removed along the axis of the strut so as to form through-openings 24, 26 without adversely impacting the strength and rigidity of the strut. Since the struts 18 and links 22 thus formed remain essentially rigid during stent expansion, the through-openings 24, 26 are also non-deforming.

[0034] The through-openings 24, 26 in the struts 18 promote the healing of the intervention site by promoting regrowth of the endothelial cells. By providing the through-openings 24, 26 in the struts, 18, the cross section of the strut is effectively reduced without decreasing the strength and integrity of the strut, as described above. As a result, the overall distance across which endothelial cell regrowth must occur is also reduced to approximately 0.0025 - 0.0035 inches, which is approximately one-half of the thickness of a convention stent. It is further believed that during insertion of the expandable medical device, cells from the endothelial layer may be scraped from the inner wall of the lumen by the through-openings 24, 26 and remain therein after implantation. The presence of such endothelial cells thus

provide a basis for the healing of the lumen.

[0035] The through-openings 24, 26 may also be loaded with an agent, most preferably a beneficial agent, for delivery to the lumen in which the tissue support device 10 is deployed.

[0036] The term "agent" as used herein is intended to have its broadest possible interpretation and is used to include any therapeutic agent or drug, as well as any body analyte, such as glucose. The terms "drug" and "therapeutic agent" are used interchangeably to refer to any therapeutically active substance that is delivered to a bodily lumen of a living being to produce a desired, usually beneficial, effect. The present invention is particularly well suited for the delivery of antiproliferatives (anti-restenosis agents) such as paclitaxil and rapamycin for example, and antithrombins such as heparin, for example. Additional uses, however, include therapeutic agents in all the major therapeutic areas including, but not limited to: anti-infectives such as antibiotics and antiviral agents; analgesics; including fentanyl, sufentanil, buprenorphine and analgesic combinations; anesthetics; anorexics; antiarthritics; antiasthmatic agents such as terbutaline; anticonvulsants; antidepressants; antidysrhythmic agents; antidiarrheals; antihistamines; anti-inflammatory agents; antimigraine preparations; anti-motion sickness preparations such as scopolamine and ondansetron; anti-nauseants; antineoplastics; antiparkinsonism drugs; antipruritics; antipsychotics; antipyretics; antispasmodics, including gastrointestinal and urinary; anticholinergics; sympathomimetics; xanthine derivatives; cardiovascular preparations, including calcium channel blockers such as nifedipine; beta blockers; beta-agonists such as dobutamine and ritodrine; antiarrhythmics; antihypertensives such as atenolol; ACE inhibitors such as ranitidine; diuretics; vasodilators, including general, coronary, peripheral, and cerebral; central nervous system stimulants; cough and cold preparations; decongestants; diagnostics; hormones such as parathyroid hormone; hypnotics; immunosuppressants; muscle relaxants; parasympatholytics; parasympathomimetics; prostaglandins; proteins; peptides; psychostimulants; sedatives; and tranquilizers.

[0037] The embodiment of the invention shown in FIGS. 1 and 2 can be further refined by using Finite Element Analysis and other techniques to optimize the deployment of the beneficial agent within the through-openings of the struts and links. Basically, the shape and location of the through-openings 24, 26 can be modified to maximize the volume of the voids while preserving the relatively high strength and rigidity of the struts 18 with respect to the ductile hinges 20.

[0038] FIG. 3 illustrates a further preferred embodiment of the present invention, wherein like reference numerals have been used to indicate like components. The tissue supporting device 100 includes a plurality of cylindrical tubes 12 connected by S-shaped bridging elements 14. Each of the cylindrical tubes 12 has a plurality of axial slots 16 extending from an end surface of the

cylindrical tube toward an opposite end surface. Formed between the slots 16 is a network of axial struts 18 and links 22. Each individual strut 18 is linked to the rest of the structure through a pair of ductile hinges 20, one at each end, which act as stress/strain concentration features. Each of the ductile hinges 20 is formed between an arc surface 28 and a concave notch surface 29. The notches 29 each have two opposed angled walls 30 which function as a stop to limit geometric deflection of the ductile hinge, and thus limit maximum device expansion.

At intervals along the neutral axis of the struts 18, at least one and more preferably a series of through-openings 24' are formed by laser drilling or any other means known to one skilled in the art. Similarly, at least one and preferably a series of through-openings 26' are formed at selected locations in the links 22. Although the use of through-openings 24' and 26' in both the struts 18 and links 22 is preferred, it should be clear to one skilled in the art that through-openings could be formed in only one of the struts and links. In the illustrated embodiment, the through-openings 24' in the struts 18 are generally rectangular whereas the through-openings 26' in the links 22 are polygonal. It should be apparent to one skilled in the art, however, that through-openings of any geometrical shape or configuration could of course be used, and that the shape of through-openings 24, 24' may be the same or different from the shape of through-openings 26, 26', without departing from the scope of the present invention. As described in detail above, the through-openings 24', 26' may be loaded with an agent, most preferably a beneficial agent, for delivery to the lumen in which the tissue support device 100 is deployed.

[0039] The relatively large, protected through-openings 24, 24', 26, 26', as described above, make the expandable medical device of the present invention particularly suitable for delivering agents having more esoteric larger molecules or genetic or cellular agents, such as, for example, protein/enzymes, antibodies, antisense, ribozymes, gene/vector constructs, and cells (including but not limited to cultures of a patient's own endothelial cells). Many of these types of agents are biodegradable or fragile, have a very short or no shelf life, must be prepared at the time of use, or cannot be preloaded into delivery devices such as stents during the manufacture thereof for some other reason. The large through-openings in the expandable device of the present invention form protected areas or receptors to facilitate the loading of such an agent at the time of use, and to protect the agent from abrasion and extrusion during delivery and implantation.

[0040] FIG. 4 shows an enlarged view of one of the struts 18 of device 100 disposed between a pair of ductile hinges 20. FIG. 5 illustrates a cross section of one of the openings 24' shown in FIG. 4. FIG. 6 illustrates the same cross section when a beneficial agent 36 has been loaded into the through-openings 24' of the struts

18. Optionally, after loading the through-openings 24' and/or the through-openings 26' with a beneficial agent 36, the entire exterior surface of the stent can be coated with a thin layer of a beneficial agent 38, which may be the same as or different from the beneficial agent 36, as schematically shown in FIG. 7. Still further, another variation of the present invention would coat the outwardly facing surfaces of the stent with a first beneficial agent 38 while coating the inwardly facing surfaces of the stent with a different beneficial agent 39, as illustrated in FIG. 8. The inwardly facing surface of the stent would be defined by at least the surfaces of the stent which, after expansion, forms the inner lumen passage. The outwardly facing surface of the stent would be defined by at least the surface of the stent which, after expansion, is in contact with and directly supports the inner wall of the lumen.

[0041] FIG. 9 illustrates yet another preferred embodiment of the present invention, wherein like reference numerals have been used to indicate like components. Unlike the stents 10, 100 described above, tissue supporting device 200 does not include through-openings which extend through the entire width of the stent. Rather, the struts 18 and/or links 22 of stent 200 preferably include at least one and preferably a plurality of recesses 40, 42, formed respectively therein on one or both side surfaces of the stent 200. The recesses 40, 42, also defined as openings, indentations, grooves, and the like, are sufficiently sized so as to promote healing of the endothelial layer and to enable a beneficial agent 36 to be loaded therein. Recesses 40, 42, like through-holes 24, 24', 26, 26', may be formed in struts 18 without compromising the strength and rigidity thereof for the same reasons as noted above. As shown above in FIGS. 7 and 8, a surface coating of one or more beneficial agents may also be provided on stent 200.

[0042] The tissue supporting device 10, 100, 200 according to the present invention may be formed of any ductile material, such as steel, gold, silver, tantalum, titanium, Nitinol, other shape memory alloys, other metals, or even some plastics. One preferred method for making the tissue supporting device 10, 100, 200 involves forming a cylindrical tube 12 and then laser cutting the slots 16, notches 29 and through-openings 24, 24', 26, 26' or recesses 40, 42 into the tube. Alternatively, the tissue supporting device may be formed by electromachining, chemical etching followed by rolling and welding, or any other method known to one skilled in the art.

[0043] The design and analysis of stress/strain concentration for ductile hinges, and stress/strain concentration features in general, is complex. The stress concentration factor can be calculated for simple ductile hinge geometries, but is generally useful only in the linear elastic range. Stress concentration patterns for a number of other geometries can be determined through photoelastic measurements and other experimental methods. Stent designs based on the use of stress/

strain concentration features, or ductile hinges, generally involve more complex hinge geometries and operate in the non-linear elastic and plastic deformation regimes.

[0044] The general nature of the relationship among applied forces, material properties, and ductile hinge geometry can be more easily understood through analysis of an idealized hinge 60 as shown in FIGS. 10a-10c. The hinge 60 is a simple beam of rectangular cross section having a width h , length L and thickness b . The idealized hinge 60 has elastic-ideally-plastic material properties which are characterized by the ideal stress/strain curve of FIG. 10d. It can be shown that the "plastic" or "ultimate bending moment" for such a beam is given by the expression:

$$M_p = M_{ult} = \delta_{yp} \frac{bh^2}{4}$$

Where b corresponds to the cylindrical tube wall thickness, h is the circumferential width of the ductile hinge, and δ_{yp} is the yield stress of the hinge material. Assuming only that expansion pressure is proportional to the plastic moment, it can be seen that the required expansion pressure to expand the tissue supporting device increases linearly with wall thickness b and as the square of ductile hinge width h . It is thus possible to compensate for relatively large changes in wall thickness b with relatively small changes in hinge width h . While the above idealized case is only approximate, empirical measurements of expansion forces for different hinge widths in several different ductile hinge geometries have confirmed the general form of this relationship. Accordingly, for different ductile hinge geometries it is possible to increase the thickness of the tissue supporting device to achieve radiopacity while compensating for the increased thickness with a much smaller decrease in hinge width.

[0045] Ideally, the stent wall thickness b should be as thin as possible while still providing good visibility on a fluoroscope. For most stent materials, including stainless steel, this would suggest a thickness of about 0.005 - 0.007 inches (0.127 - 0.178 mm) or greater. The inclusion of ductile hinges in a stent design can lower expansion forces/pressures to very low levels for any material thickness of interest. Thus ductile hinges allow the construction of optimal wall thickness tissue supporting devices at expansion force levels significantly lower than current non-visible designs.

[0046] The expansion forces required to expand the tissue supporting device 10, 100, 200 according to the present invention from an initial condition illustrated in FIG. 1 to an expanded condition is between 1 and 5 atmospheres, preferably between 2 and 3 atmospheres. The expansion may be performed in a known manner, such as by inflation of a balloon or by a mandrel. The tissue supporting device 10, 100, 200 in the expanded

condition has a diameter which is preferably up to three times the diameter of the device in the initial unexpanded condition.

[0047] Many tissue supporting devices fashioned from cylindrical tubes comprise networks of long, narrow, prismatic beams of essentially rectangular cross section as shown in FIG. 11. These beams which make up the known tissue supporting devices may be straight or curved, depending on the particular design. Known expandable tissue supporting devices have a typical wall thickness b of 0.0025 inches (0.0635 mm), and a typical strut width h of 0.005 to 0.006 inches (0.127 - 0.1524 mm). The ratio of $b:h$ for most known designs is 1:2 or lower. As b decreases and as the beam length L increases, the beam is increasingly likely to respond to an applied bending moment M by buckling, and many designs of the prior art have displayed this behavior. This can be seen in the following expression for the "critical buckling moment" for the beam of FIG. 6.

$$M_{crit} = \frac{\pi b^3 h \sqrt{EG(1-0.63 b/h)}}{6L}$$

Where:

E = Modulus of Elasticity

G = Shear Modulus

[0048] By contrast, in a ductile hinge based design according to the present invention, only the hinge itself deforms during expansion. The typical ductile hinge 20 is not a long narrow beam as are the struts in the known stents. Wall thickness of the present invention may be increased to 0.005 inches (0.127 mm) or greater, while hinge width is typically 0.002 - 0.003 inches (0.0508 - 0.0762 mm), preferably 0.0025 inches (0.0635 mm) or less. Typical hinge length, at 0.002 to 0.005 inches (0.0508 - 0.127 mm), is more than an order of magnitude less than typical strut length. Thus, the ratio of $b:h$ in a typical ductile hinge 20 is 2:1 or greater. This is an inherently stable ratio, meaning that the plastic moment for such a ductile hinge beam is much lower than the critical buckling moment M_{crit} , and the ductile hinge beam deforms through normal strain-curvature. Ductile hinges 20 are thus not vulnerable to buckling when subjected to bending moments during expansion of the tissue supporting device 10, 100, 200.

[0049] To provide optimal recoil and crush-strength properties, it is desirable to design the ductile hinges so that relatively large strains, and thus large curvatures, are imparted to the hinge during expansion of the tissue supporting device. Curvature is defined as the reciprocal of the radius of curvature of the neutral axis of a beam in pure bending. A larger curvature during expansion results in the elastic curvature of the hinge being a small fraction of the total hinge curvature. Thus, the gross elastic recoil of the tissue supporting device is a

small fraction of the total change in circumference. It is generally possible to do this because common stent materials, such as 316L Stainless Steel have very large elongations-to-failure (i.e., they are very ductile).

[0050] It is not practical to derive exact expressions for residual curvatures for complex hinge geometries and real materials (i.e., materials with non-idealized stress/strain curves). The general nature of residual curvatures and recoil of a ductile hinge may be understood by examining the moment-curvature relationship for the elastic-ideally-plastic rectangular hinge 60 shown in FIGS. 10a-c. It may be shown that the relationship between the applied moment and the resulting beam curvature is:

$$M = M_p [1 - \frac{1}{3} (\frac{\kappa_0}{\kappa_{yp}})^2] = \frac{3}{2} M_{yp} [1 - \frac{1}{3} (\frac{\kappa_{yp}}{\kappa})^2]$$

This function is plotted in FIG. 12. It may be seen in this plot that the applied moment M asymptotically approaches a limiting value M_p , called the plastic or ultimate moment. Beyond $\frac{11}{12} M_p$ large plastic deformations occur with little additional increase in applied moment. When the applied moment is removed, the beam rebounds elastically along a line such as a-b. Thus, the elastic portion of the total curvature approaches a limit of $\frac{3}{2}$ the curvature at the yield point. These relations may be expressed as follows:

$$M_p = \frac{3}{2} M_{yp} \Rightarrow \kappa_{rebound} = \frac{3}{2} \kappa_{yp}$$

[0051] Imparting additional curvature in the plastic zone cannot further increase the elastic curvature, but will decrease the ratio of elastic to plastic curvature. Thus, additional curvature or larger expansion of the tissue supporting device will reduce the percentage recoil of the overall stent structure.

[0052] As shown in FIG. 13, when a rigid strut 18 is linked to the ductile hinge 60 described above, the strut 18 forms an angle θ with the horizontal that is a function of hinge curvature. A change in hinge curvature results in a corresponding change in this angle θ . The angular elastic rebound of the hinge is the change in angle $\Delta\theta$ that results from the rebound in elastic curvature described above, and thus angular rebound also approaches a limiting value as plastic deformation proceeds. The following expression gives the limiting value of angular elastic rebound for the idealized hinge of FIG. 13.

$$\theta_{rebound} = 3\epsilon_{yp} \frac{L}{h}$$

Where strain at the yield point is an independent material property (yield stress divided by elastic modulus); L is the length of the ductile hinge; and h is the width of

the hinge. For non-idealized ductile hinges made of real materials, the constant 3 in the above expression is replaced by a slowly rising function of total strain, but the effect of geometry would remain the same. Specifically, the elastic rebound angle of a ductile hinge decreases as the hinge width h increases, and increases as the hinge length L increases. To minimize recoil, therefore, hinge width h should be increased and length L should be decreased.

[0053] Ductile hinge width h will generally be determined by expansion force criteria, so it is important to reduce hinge length to a practical minimum in order to minimize elastic rebound. Empirical data on recoil for ductile hinges of different lengths show significantly lower recoil for shorter hinge lengths, in good agreement with the above analysis.

[0054] The ductile hinges 20 of the tissue supporting device 10, 100, 200 provide a second important advantage in minimizing device recoil. The embodiment of FIG. 1 shows a network of struts joined together through ductile hinges to form a cylinder. As the device is expanded, curvature is imparted to the hinges 20, and the struts 18 assume an angle θ with respect to their original orientation, as shown in FIG. 13. The total circumferential expansion of the tissue supporting device structure is a function of hinge curvature (strut angle) and strut length. Moreover, the incremental contribution to stent expansion (or recoil) for an individual strut depends on the instantaneous strut angle. Specifically, for an incremental change in strut angle $\Delta\theta$, the incremental change in circumference ΔC will depend on the strut length R and the cosine of the strut angle θ .

$$\Delta C = R \Delta\theta \cos\theta$$

[0055] Since elastic rebound of hinge curvature is nearly constant at any gross curvature, the net contribution to circumferential recoil ΔC is lower at higher strut angles θ . The final device circumference is usually specified as some fixed value, so decreasing overall strut length can increase the final strut angle θ . Total stent recoil can thus be minimized with ductile hinges by using shorter struts and higher hinge curvatures when expanded.

[0056] Empirical measurements have shown that tissue supporting device designs based on ductile hinges, such as the embodiment of FIG. 1, display superior resistance to compressive forces once expanded despite their very low expansion force. This asymmetry between compressive and expansion forces may be due to a combination of factors including the geometry of the ductile hinge, the increased wall thickness, and increased work hardening due to higher strain levels.

[0057] According to one example of the tissue supporting device of the invention, the device can be expanded by application of an internal pressure of about 2 atmospheres or less, and once expanded to a diam-

eter between 2 and 3 times the initial diameter can withstand a compressive force of about 16 to 20 gm/mm or greater. Examples of typical compression force values for prior art devices are 3.8 to 4.0 gm/mm.

[0058] While both recoil and crush strength properties of tissue supporting devices can be improved by use of ductile hinges with large curvatures in the expanded configuration, care must be taken not to exceed an acceptable maximum strain level for the material being used. Generally, ϵ_{max} is defined as maximum strain, and it is dependent on ductile hinge width h , ductile hinge length L , and bend angle θ in radians. When strain, hinge width and bend angle are determined through other criteria, an expression may be developed to determine the required lengths for the complicated ductile hinge geometry of the present invention. Typical values for the prismatic portions of the curved ductile hinges 20 range from about 0.002 to about 0.0035 inches (0.051 - 0.089 mm) in hinge width and about 0.002 to about 0.006 inches (0.051 - 0.152 mm) in hinge length.

[0059] In many designs of the prior art, circumferential expansion was accompanied by a significant contraction of the axial length of the stent which may be up to 15% of the initial device length. Excessive axial contraction can cause a number of problems in device deployment and performance including difficulty in proper placement and tissue damage. Designs based on ductile hinges 20 can minimize the axial contraction, or foreshortening, of a tissue supporting device during expansion, as discussed in greater detail in the aforementioned U.S. Application Serial No. 09/183,555. This ability to control axial contraction based on hinge and strut design provides great design flexibility when using ductile hinges. For example, a stent could be designed with zero axial contraction.

[0060] The stent 10, 100, 200 of the present invention illustrates the trade off between crush strength and axial contraction. Referring to FIG. 3, a portion of the tissue supporting device 100 having an array of struts 18 and ductile hinges 20 are shown in the unexpanded state. The struts 18 are positioned initially at an angle θ_1 with respect to a longitudinal axis X of the device. As the device is expanded radially from the unexpanded state illustrated in FIG. 3, the angle θ_1 increases. In this case the device contracts axially from the onset of vertical expansion throughout the expansion. A higher final strut angle θ_1 can significantly increase crush strength and decrease circumferential recoil of the stent structure. However, there is a trade off between increased crush strength and increase in axial contraction.

[0061] According to one example of the present invention, the struts 18 are positioned initially at an angle of about 0° to 45° with respect to a longitudinal axis of the device. As the device is expanded radially from the unexpanded state illustrated in FIG. 3, the strut angle increases to about 20° to 80°.

[0062] In addition, while ductile hinges 20 are the preferred configuration for the expandable medical device

of the present invention, a stent without the defined ductile hinges would also be included within the scope of the present invention.

[0063] While the invention has been described in detail with reference to the preferred embodiments thereof, it will be apparent to one skilled in the art that various changes and modifications can be made and equivalents employed, without departing from the present invention.

The expandable medical device may comprise: at least one cylindrical tube; a network of elongated struts formed in said cylindrical tube, wherein each of the elongated struts are axially displaced from adjacent struts; and a plurality of ductile hinges formed between the elongated struts, the ductile hinges allowing the cylindrical tube to be expanded or compressed from a first diameter to a second diameter by deformation of the ductile hinges; wherein at least one of the elongated struts includes at least one opening for loading of a beneficial agent therein. The expandable medical device may comprise a plurality of openings. The expandable medical device may comprise at least one opening extending through a thickness of said at least one strut, so as to thereby define a through-opening. The expandable medical device may also comprise at least one opening having a depth less than a thickness of said at least one strut, so as to thereby define a recess. The expandable medical device may further comprise a beneficial agent loaded within said at least one opening; said beneficial agent may include antiproliferatives or antithrombins. Said beneficial agent may also include at least one of large molecules, microspheres, beneficial agents within a biodegradable polymer carrier, and cells, said at least one opening of said at least one strut forming a protected receptor for loading said beneficial agent therein. The expandable medical device may further comprise a surface coating of a second beneficial agent on said expandable medical device. Said first and second beneficial agents may be different, or identical. The surface coating of said second beneficial agent may cover a portion of said expandable medical device and may further comprise a surface coating of a third beneficial agent covering another portion of said expandable medical device, wherein said second and third beneficial agents are different. The expandable medical device may include two cylindrical tubes interconnected by a bridging element. A transition between the cross-sectional area of the struts and the cross-sectional area of the ductile hinges may be an abrupt transition which extends less than 10 percent of a length of a strut. The expandable medical devices may have a plurality of ductile hinges which may be curved prismatic beams having an arc surface and a concave surface. The expandable medical device may further comprise a geometric deflection limiting feature for limiting an amount of bending of the ductile hinges. The deflection limiting feature may include angled side walls on opposite sides of a ductile hinge which engage one another to limit deflection at

ductile hinge. The deflection limiting members may be V-shaped notches. The ductile hinges may be designed to deform plastically upon radial expansion or compression of the expandable medical device while the elongated struts experience no plastic deformation upon radial expansion or compression. The expandable medical device may be formed of Nitinol, or of stainless steel. The expandable medical device may also comprise a plurality of elongated struts, said plurality of elongated struts joined together to form a substantially cylindrical device which is expandable from a cylinder having a first diameter to a cylinder having a second diameter, said plurality of struts each having a strut width in a circumferential direction and a strut thickness in a radial direction; and wherein at least one of said plurality of struts includes at least one recess extending at least partially through a thickness of said strut. Said at least one recess may extend entirely through the thickness of said strut so as to define a through-opening. Said at least one recess may be generally rectangular, or polygonal.

Claims

1. An expandable medical device (100), comprising:

a plurality of elongated struts (18) forming a substantially cylindrical device (12) which is expandable from a cylinder having a first diameter to a cylinder having a second diameter;

hinges (20) formed at said elongated struts (18) to substantially prevent deformation of said elongated struts (18) during expansion of said cylindrical device (12), and

links (22) arranged between said elongated struts (18);

wherein a series of openings (24, 26; 24', 26') are formed in said struts (18) and said links (22).

2. An expandable medical device (100), comprising:

a plurality of elongated struts (18) forming a substantially cylindrical device (12) which is expandable from a cylinder having a first diameter to a cylinder having a second diameter, and

hinges (20) formed at said elongated struts (18) to substantially prevent deformation of said elongated struts (18) during expansion of said cylindrical device (12);

wherein a series of openings (24; 24') is formed only in said struts (18).

3. An expandable medical device (100) comprising:

a plurality of elongated struts (18) forming a substantially cylindrical device (12) which is expandable from a cylinder having a first diameter to a cylinder having a second diameter;

hinges (20) formed at said elongated struts (18) to substantially prevent deformation of said elongated struts (18) during expansion of said cylindrical device (12), and

links (22) arranged between said elongated struts (18),

wherein a series of openings (26; 26') is formed only in said links (22).

4. The expandable medical device according to one of the preceding claims, wherein said series of openings extends entirely through the thickness of said strut so as to define through-openings.

5. The expandable medical device according to one of the preceding claims, wherein at least one of said openings is generally rectangular.

6. The expandable medical device according to one of the preceding claims, wherein at least one of said openings is polygonal.

7. The expandable medical device according to one of the preceding claims, further comprising a beneficial agent loaded within at least one of said openings.

8. The expandable medical device according to Claim 7, wherein the beneficial agent undergoes substantially no deformation when said expandable medical device is expanded from the first diameter to the second diameter.

9. The expandable medical device according to Claim 7 or 8, wherein said beneficial agent comprises a plurality of beneficial agents.

10. The expandable medical device according to one of Claims 7 to 9, wherein said beneficial agent includes at least one antiproliferative.

11. The expandable medical device according to Claim 7 or 8, wherein said beneficial agent is paclitaxel.

12. The expandable medical device according to Claim 7 to 10, wherein said beneficial agent is rapamycin.

13. The expandable medical device according to one of Claims 7 to 10, wherein said beneficial agent includes antithrombins.

14. The expandable medical device according to one of Claims 7 to 10, or Claim 13, wherein said beneficial agent includes at least one of large molecules, microspheres, beneficial agents within a biodegradable polymer carrier, and cells, said at least one opening forming a protected receptor for loading said beneficial agent therein. 5
15. The expandable medical device according to one of the preceding claims, wherein the hinges formed between the elongated struts comprise a plurality of ductile hinges providing means for concentration of stress/strain, the ductile hinges allowing the cylindrical tube to be expanded or compressed from the first diameter to the second diameter by deformation of the ductile hinges. 10
16. The expandable medical device according to Claim 15, wherein a transition between the cross-sectional area of the struts and the cross-sectional area of the ductile hinges is an abrupt transition which extends less than 10 percent of a length of a strut. 20
17. The expandable medical device according to Claim 15 or 16, wherein the plurality of ductile hinges are curved prismatic beams having an arc surface and a concave surface. 25
18. The expandable medical device according to one of Claims 15 to 17, wherein the ductile hinges are designed to deform plastically upon radial expansion or compression of the expandable medical device while the elongated struts experience no plastic deformation upon radial expansion or compression. 30 35
19. A method of forming an expandable medical device, comprising:
- cutting a plurality of slots in a substantially cylindrical tube to form a plurality of elongated struts forming a substantially cylindrical device which is expandable from a cylinder having a first diameter to a cylinder having a second diameter; and 40 45
- creating means for concentration of stress/strain between said elongated struts to substantially prevent deformation of said elongated struts during expansion of the cylinder; 50
- characterized in that
- a plurality of recesses are drilled in the elongated struts such that the plurality of recesses are non-deforming when said cylinder is expanded from the first diameter to the second diameter. 55
20. The method according to Claim 19, wherein the plurality of slots are formed by laser cutting.
21. The method according to Claim 19 or 20, wherein the plurality of recesses are formed by laser drilling.
22. The method according to one of Claims 19 to 21, further comprising filling the plurality of recesses with a beneficial agent.
23. The method according to Claim 22, wherein said beneficial agent includes at least one antiproliferative.
24. The method according to Claim 22, wherein said beneficial agent is paclitaxel.
25. The method according to Claim 22, wherein said beneficial agent is rapamycin.

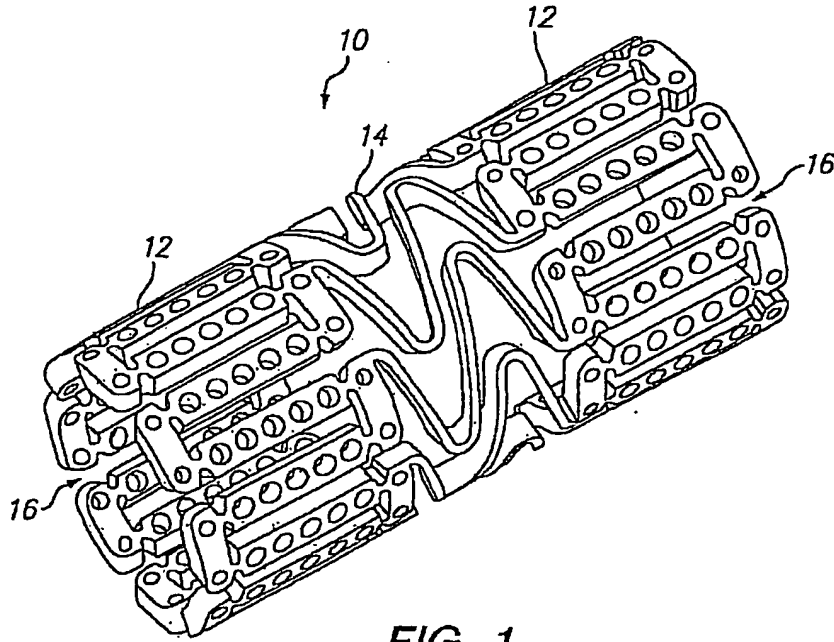


FIG. 1

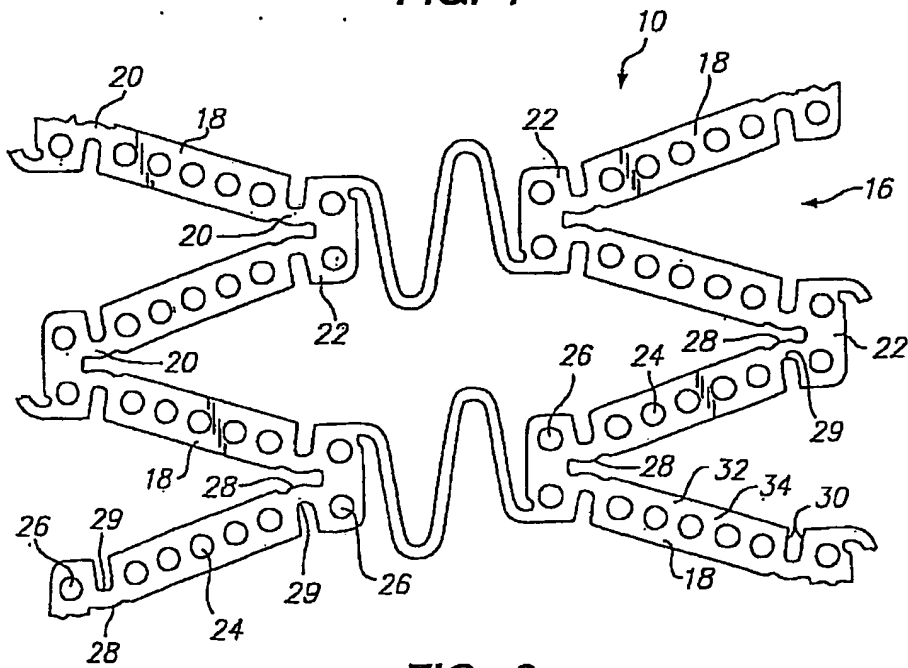


FIG. 2

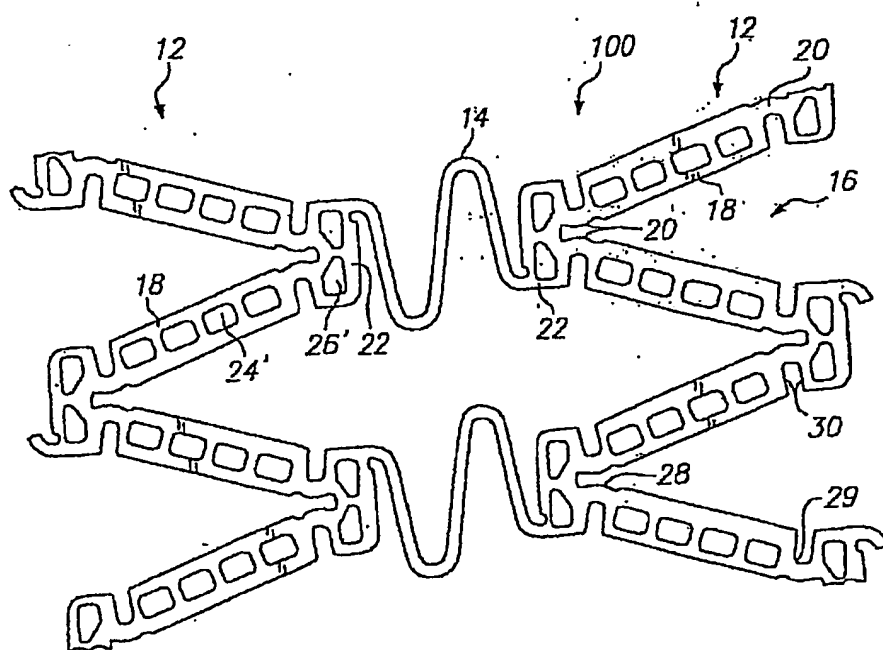
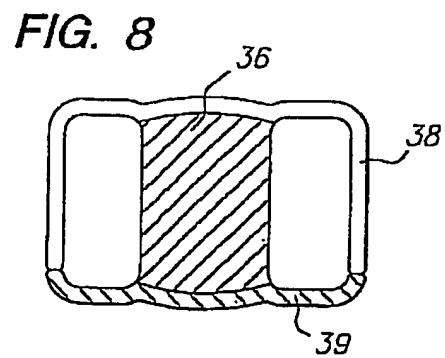
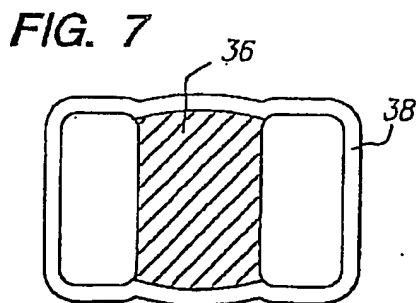
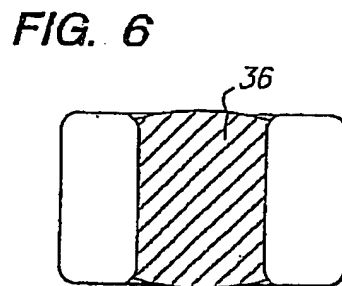
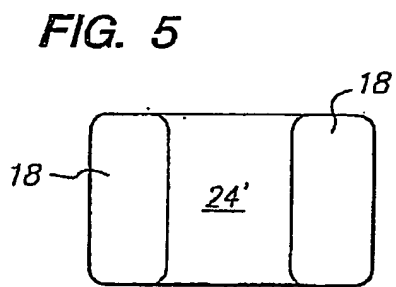
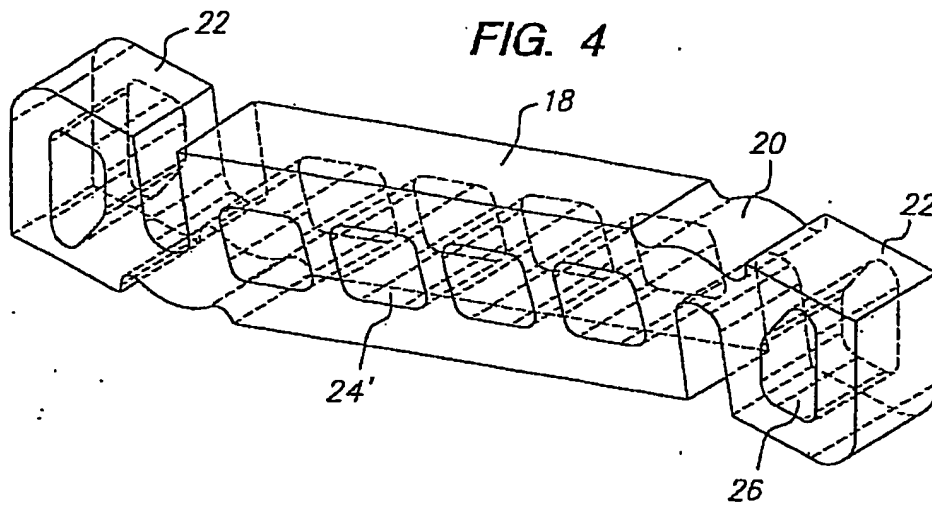
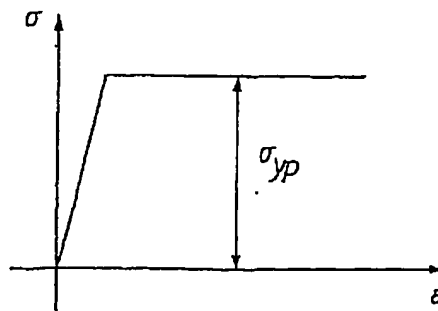
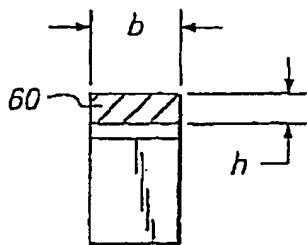
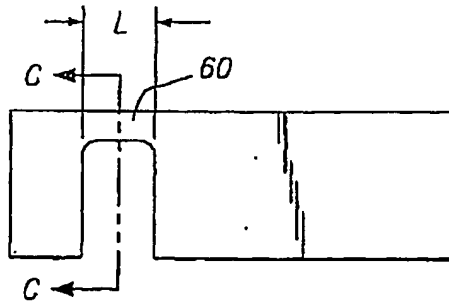
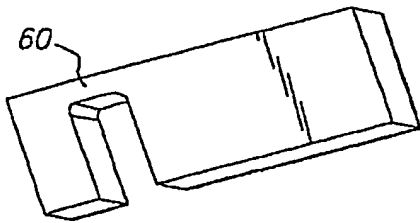
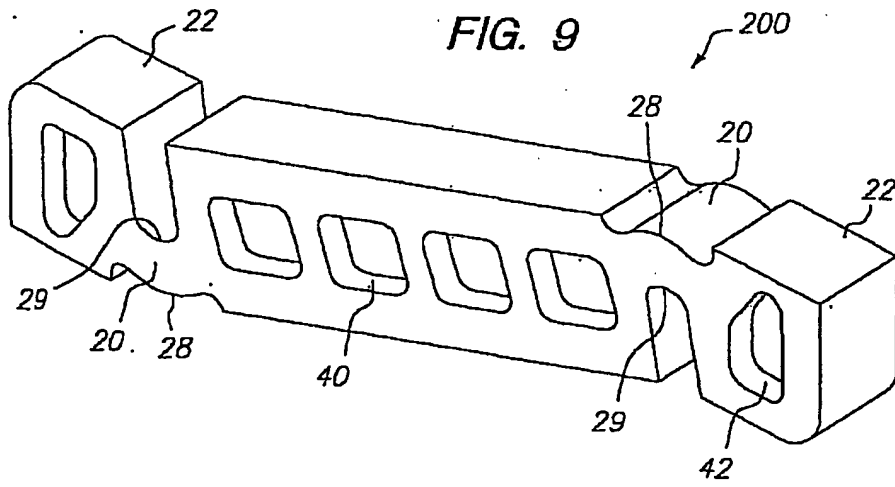
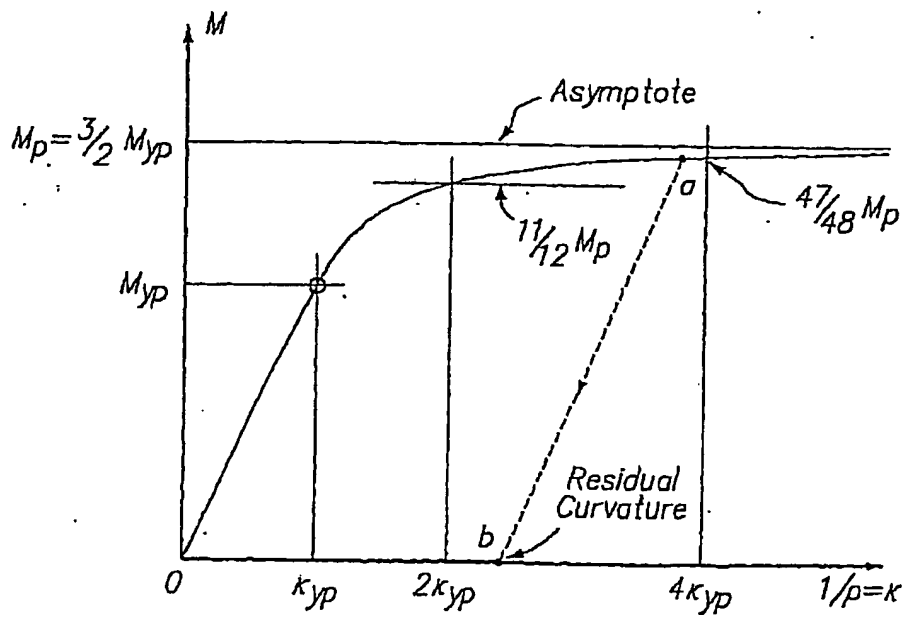
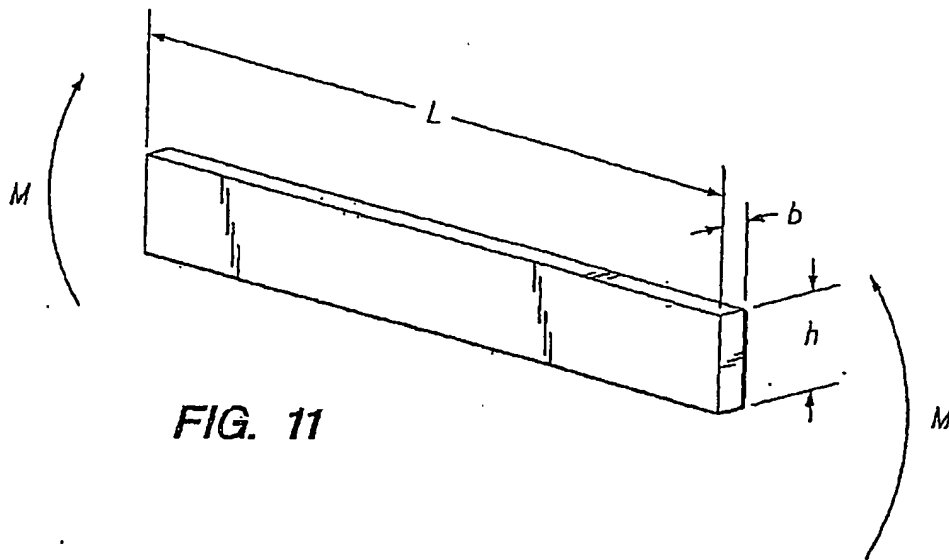


FIG. 3







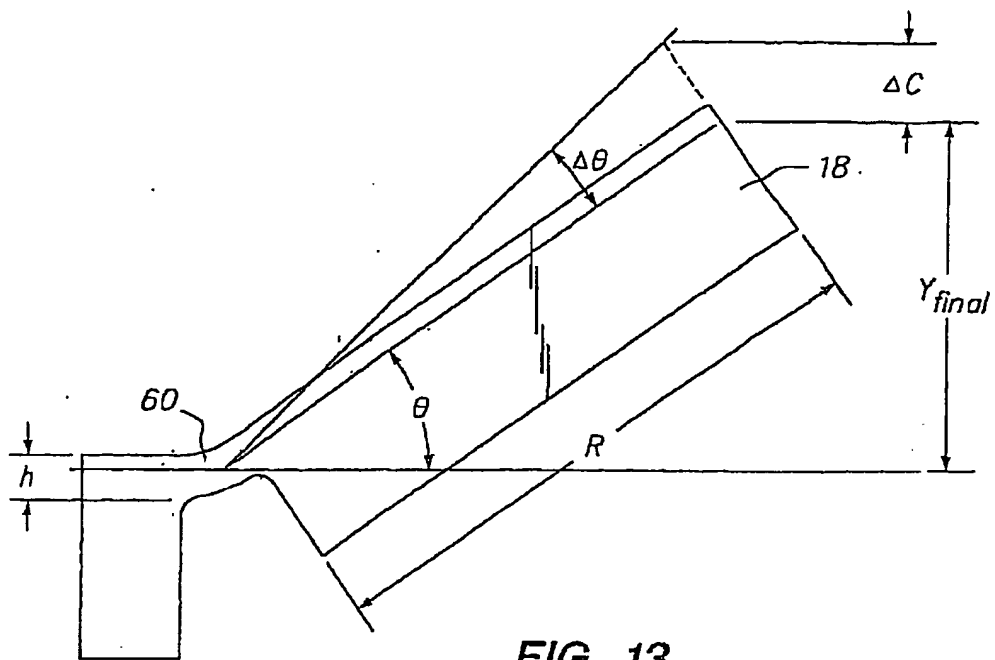


FIG. 13



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 05 01 3944

| DOCUMENTS CONSIDERED TO BE RELEVANT | | | |
|--|--|--|--|
| Category | Citation of document with indication, where appropriate, of relevant passages | Relevant to claim | CLASSIFICATION OF THE APPLICATION (In I.C.I.7) |
| X | EP 0 950 386 A (CORDIS CORP) 20 October 1999 (1999-10-20) | 1-3,5-8, 10-12, 15-18 9,13,14 4,19-25 | A61F2/06 |
| Y | * paragraph [0034] * | | |
| A | * figure 2A * | | |
| | * column 8, line 2 - line 12 * | | |
| | ----- | | |
| X | US 5 972 027 A (JOHNSON MICHAEL W) 26 October 1999 (1999-10-26) | 1 | |
| Y | * figures 2,5 * | 9 | |
| | ----- | | |
| Y | US 6 096 070 A (BATES BRIAN L ET AL) 1 August 2000 (2000-08-01) | 13,14 | |
| | * column 7, line 55 - column 8, line 13 * | | |
| | * column 9, line 20 - column 10, line 11 * | | |
| | * column 16, line 32 - column 17, line 4 * | | |
| | * column 20, line 6 - line 14 * | | |
| | * figures 8-10D * | | |
| | ----- | | |
| A | WO 99/49928 A (SHANLEY JOHN F) 7 October 1999 (1999-10-07) | 1-25 | |
| | * page 13, last paragraph - page 15, paragraph 1 * | | |
| | * page 24, last paragraph - page 25, paragraph 1 * | | |
| | * figures 3E,4A * | | |
| | ----- | | |
| A | WO 00/45744 A (SCIMED LIFE SYSTEMS INC ;WANG LIXIAO (US); STANSLASKI JOEL (US); Y) 10 August 2000 (2000-08-10) | 1-25 | |
| | * page 4, line 17 - page 6, line 5 * | | |
| | * figure 3 * | | |
| | ----- | | |
| | -/-- | | |
| The present search report has been drawn up for all claims | | | |
| Place of search Munich | | Date of completion of the search 10 August 2005 | Examiner Franz, V |
| <p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons</p> <p>& : member of the same patent family, corresponding document</p> | | | |

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EPO FORM 1503 (02.02.99) (P4/C01)



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 05 01 3944

| DOCUMENTS CONSIDERED TO BE RELEVANT | | | |
|--|---|--|--|
| Category | Citation of document with indication, where appropriate, of relevant passages | Relevant to claim | CLASSIFICATION OF THE APPLICATION (In I.C.I.7) |
| A | WO 96/32907 A (SCHNEIDER USA INC) 24 October 1996 (1996-10-24) * page 3, line 28 - line 31 * * page 4, line 23 - page 5, line 7 * ----- | 1-25 | |
| | | | TECHNICAL FIELDS SEARCHED (In I.C.I.7) |
| | | | |
| The present search report has been drawn up for all claims | | | |
| Place of search Munich | | Date of completion of the search 10 August 2005 | Examiner Franz, V |
| CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document | | T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document | |

3
EPO FORM 1503 (3.12.02) (POL/C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 05 01 3944

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

10-08-2005

| Patent document cited in search report | | Publication date | Patent family member(s) | Publication date |
|---|---|---------------------|----------------------------|---------------------|
| EP 0950386 | A | 20-10-1999 | US 6273913 B1 | 14-08-2001 |
| | | | AT 263520 T | 15-04-2004 |
| | | | CA 2269310 A1 | 16-10-1999 |
| | | | DE 69916157 D1 | 13-05-2004 |
| | | | DE 69916157 T2 | 17-03-2005 |
| | | | EP 1407726 A1 | 14-04-2004 |
| | | | EP 0950386 A2 | 20-10-1999 |
| | | | ES 2218953 T3 | 16-11-2004 |
| | | | PT 950386 T | 31-08-2004 |
| | | | US 2003176915 A1 | 18-09-2003 |
| | | | US 2001029351 A1 | 11-10-2001 |
| | | | US 2001027340 A1 | 04-10-2001 |
| | | | US 2005085902 A1 | 21-04-2005 |
| | | | | |
| US 5972027 | A | 26-10-1999 | WO 9916386 A1 | 08-04-1999 |
| | | | US 6253443 B1 | 03-07-2001 |
| | | | US 2001029660 A1 | 18-10-2001 |
| | | | | |
| US 6096070 | A | 01-08-2000 | US 5609629 A | 11-03-1997 |
| | | | AU 716005 B2 | 17-02-2000 |
| | | | AU 5588896 A | 19-12-1996 |
| | | | CA 2178541 A1 | 08-12-1996 |
| | | | US 6774278 B1 | 10-08-2004 |
| | | | WO 9817331 A1 | 30-04-1998 |
| | | | US 2003028243 A1 | 06-02-2003 |
| | | | US 2003028244 A1 | 06-02-2003 |
| | | | US 2003036794 A1 | 20-02-2003 |
| | | | US 2004047909 A1 | 11-03-2004 |
| | | | US 5824049 A | 20-10-1998 |
| | | | US 5873904 A | 23-02-1999 |
| | | | US 2004243225 A1 | 02-12-2004 |
| | | | DE 69623855 D1 | 31-10-2002 |
| | | | DE 69623855 T2 | 28-05-2003 |
| | | | DK 747069 T3 | 09-12-2002 |
| | | | EP 0747069 A2 | 11-12-1996 |
| | | | ES 2184838 T3 | 16-04-2003 |
| | | | JP 9099056 A | 15-04-1997 |
| | | | | |
| | | | | |
| WO 9949928 | A | 07-10-1999 | US 6241762 B1 | 05-06-2001 |
| | | | AU 773731 B2 | 03-06-2004 |
| | | | AU 3208799 A | 18-10-1999 |
| | | | AU 2004202044 A1 | 10-06-2004 |
| | | | CA 2323358 A1 | 07-10-1999 |
| | | | EP 1222941 A2 | 17-07-2002 |
| | | | EP 1071490 A1 | 31-01-2001 |
| | | | JP 2002509775 T | 02-04-2002 |

EPD FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

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10-08-2005

| Patent document cited in search report | | Publication date | Patent family member(s) | Publication date |
|---|---|---------------------|----------------------------|---------------------|
| WO 9949928 | A | | JP 2005034670 A | 10-02-2005 |
| | | | US 2005159806 A1 | 21-07-2005 |
| | | | WO 9949928 A1 | 07-10-1999 |
| | | | US 2003009214 A1 | 09-01-2003 |
| | | | US 2003167085 A1 | 04-09-2003 |
| | | | US 2003199970 A1 | 23-10-2003 |
| | | | US 6562065 B1 | 13-05-2003 |
| | | | US 2004122505 A1 | 24-06-2004 |
| | | | US 2004193249 A1 | 30-09-2004 |
| | | | US 2004254635 A1 | 16-12-2004 |
| | | | US 2004225350 A1 | 11-11-2004 |
| | | | US 2004236408 A1 | 25-11-2004 |
| | | | US 6293967 B1 | 25-09-2001 |
| | | | US 2002165604 A1 | 07-11-2002 |
| | | | US 2002013619 A1 | 31-01-2002 |
| WO 0045744 | A | 10-08-2000 | US 6419692 B1 | 16-07-2002 |
| | | | AU 2724800 A | 25-08-2000 |
| | | | EP 1150622 A1 | 07-11-2001 |
| | | | JP 2002536058 T | 29-10-2002 |
| | | | WO 0045744 A1 | 10-08-2000 |
| | | | US 2002151844 A1 | 17-10-2002 |
| WO 9632907 | A | 24-10-1996 | AT 223180 T | 15-09-2002 |
| | | | AU 4952096 A | 07-11-1996 |
| | | | BR 9608021 A | 02-03-1999 |
| | | | CA 2216943 A1 | 24-10-1996 |
| | | | DE 29624503 U1 | 16-09-2004 |
| | | | DE 69623455 D1 | 10-10-2002 |
| | | | DE 69623455 T2 | 16-01-2003 |
| | | | EP 0822788 A1 | 11-02-1998 |
| | | | WO 9632907 A1 | 24-10-1996 |
| | | | JP 10506560 T | 30-06-1998 |
| | | | NO 974823 A | 17-10-1997 |
| | | | US 6358556 B1 | 19-03-2002 |
| | | | US 2002091433 A1 | 11-07-2002 |
| | | | US 2002071902 A1 | 13-06-2002 |
| | | | US 2002032477 A1 | 14-03-2002 |
| | | | US 2004049265 A1 | 11-03-2004 |
| | | | US 5837313 A | 17-11-1998 |
| | | | US 6120536 A | 19-09-2000 |
| | | | US 2002004101 A1 | 10-01-2002 |
| | | | US 2005106210 A1 | 19-05-2005 |
| | | | ZA 9603075 A | 20-10-1997 |

EPO FORM P4559

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

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